- 2. The method of claim 1, wherein said patient is administered a pharmaceutical composition comprising said noribogaine and wherein said noribogaine is the sole analgesic agent in said pharmaceutical composition.
- 3. A method of alleviating pain in a patient for whom opioid analgesics are contraindicated, comprising: administering systemically an amount of noribogaine to said patient effective to reduce or eliminate pain in said patient in the absence of any concomitant opioid analgesic therapy.
- 4. The method of any of claims 1-3, wherein said noribogaine is administered to said patient at a dose of between 0.1 mg and 100 mg per kg of body weight.
- 5. The method of claim 4, wherein said noribogaine is administered at a dose of between 1.0 mg and 30 mg per kg of body weight.
- 6. A method of treating a patient to alleviate pain, comprising:
 - a) administering systemically to said patient an amount of noribogaine; and
 - b) concomitantly administering systemically to said patient an amount of one or more opioid antagonists;

wherein said respective amounts of noribogaine and said one or more opioid antagonists are effective to reduce or eliminate pain in said patient.

- 7. The method of claim 6, wherein said opioid antagonist is naloxone, administered to said patient at a dose between 0.05 mg and 0.5 mg for each mg of noribogaine.
- 8. The method of claim 6, wherein said opioid antagonist is naltrexone, administered to said patient at a dose of between 0.05 mg and 0.5 mg for each mg of noribogaine.
- 9. The method of claim 6, wherein said noribogaine and said opioid antagonist are administered transdermally.

Remarks

After amendment, claims 1-9 remain pending in the present application, claims 10-24